Federal Regulations, Section 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: May 18, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95–12779 Filed 5–24–95; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 20, 1994, and published in the **Federal Register** on October 28, 1994, (59 FR 54219), Hoffman-LaRoche, Inc., 340 Kingsland Street, Nutley, New Jersey 07110, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Levorphanol (9220), a basic class of controlled substance listed in Schedule II.

No comments or objections have been received. Therefore, pursuant to Section 303 of the Comprehensive Drug Abuse Prevention and Control Act of 1970 and Title 21, Code of Federal Regulations, Section 1301.54(e), the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: May 17, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95–12780 Filed 5–24–95; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Correction

In the **Federal Register** (FR Doc. 95–8920) Vol. 60, No. 70 at page 18618, dated April 12, 1995, the listing of controlled substances should have included Oxycodone (9143), Hydromorphone (9150), Diphenoxylate (9170) and Noroxymorphone (9668) for Mallinckrodt Chemical, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147.

Dated: May 17, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95–12778 Filed 5–24–95; 8:45 am]

BILLING CODE 4410–09–M

Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on January 18, 1995, North Pacific Trading Company, 1505 SE Gideon Street, Portland, Oregon 97202, made application to the Drug Enforcement Administration to be registered as an importer of Marihuana (7360) a basic class of controlled substance in Schedule I.

This application is exclusively for the importation of marithuana seed which will be rendered non-viable and used as bird seed.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.54 in such form as prescribed by 21 CFR 1305.47.

Any such comments, objections, or requests for a hearing may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justices, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 26, 1995.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43747–46 (September 23, 1975), all applicants for registration to import basic classes of any controlled substances in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1311. 42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: May 18, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95–12783 Filed 5–24–95; 8:45 am]

BILLING CODE 4410–09–M

Importer of Controlled Substances; Application Withdrawal for Nycomed Incorporated

By letter dated April 17, 1995, Nycomed Inc., 33 Riverside Avenue, Rensselaer, New York 12144, withdrew their request to be registered as an importer of Meperidine (9230).

Therefore, the Notice dated February 14, 1995, in **Federal Register** (FR Doc. 95–3627), Vol. 60, No. 30 at page 8414 is hereby withdrawn.

Dated: May 17, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95–12781 Filed 5–24–95; 8:45 am]

BILLING CODE 4410–09–M

Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on March 22, 1995, Research Biochemicals, Limited Partnership, One Strathmore Road, Natick, Massachusetts 01760, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below;

Drug	Schedule
Methaqualone (2565)	I
Ibogaine (7260)	I
Tetrahydrocannabinois (7370)	I
Bufotenie (7433)	I
Dimethyltryptamine (7435)	I
Etorphine (except HC1) (9056)	I
Methylphenidate (1724)	II
Etorphine Hydrochloride (9059)	II
Diphenoxylate (9170)	II

Drug	Schedule
Metazocine (9240)	II
Methadone (9250)	II
Fentanyl (9801)	II

The firm plans to import small quantities of the controlled substances to manufacture laboratory reference standards and neurochemicals.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comment son or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.54 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 26, 1995.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42 (b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import basic classes of any controlled substances in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: May 17, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95–12784 Filed 5–24–95; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 22, 1995, Roche Diagnostic Systems Inc., 1080 U.S. Highway 202, Somerville, New Jersey 08876, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Phencyclidine (7471)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to manufacture very small quantities of the listed controlled substances which will be incorporated in drug of abuse detection kits.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the above application and may also file a written request for a hearing thereon in accordance with 21 CFR 1301.54 and in the form prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 26, 1995.

Dated May 18, 1995.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 95–12785 Filed 5–24–95; 8:45 am]

BILLING CODE 4410–09–M

Immigration and Naturalization Service

[INS No. 1723–95; AG Order No. 1967–95]

RIN 1115–AC30

Extension of Designation of Rwanda Under Temporary Protected Status Program

AGENCY: Immigration and Naturalization Service, Justice.

ACTION: Notice.

SUMMARY: This notice extends, until June 6, 1996, the Attorney General's designation of Rwanda under the Temporary Protected Status program provided for in section 244A of the Immigration and Nationality Act, as amended (''the Act''). Accordingly, eligible aliens who are nationals of Rwanda, or who have no nationality and who last habitually resided in Rwanda, may re-register for Temporary Protected Status and extension of employment authorization. This re-registration is limited to persons who already have registered or will register for the initial period of Temporary Protected Status, which ends on June 6, 1995. In addition during the extension period, some aliens may be eligible for late initial registration pursuant to 8 CFR 240.2(f)(2).

EFFECTIVE DATE: This extension of designation is effective on June 7, 1995, and will remain in effect until June 6, 1996. Re-registration procedures become effective May 25, 1995, and will remain in effect until June 23, 1995.

FOR FURTHER INFORMATION CONTACT: Ronald Chirlin, Senior Immigration Examiner, Immigration and Naturalization Service, Room 3214, 325 I Street, NW., Washington, DC 20536, telephone (202) 514–5014.

SUPPLEMENTARY INFORMATION: Under section 244A of the Act, as amended by section 302(a) of Pub. L. 101–649 and section 304(b) of Pub. L. 102–232 (8 U.S.C. 1254a), the Attorney General is authorized to grant Temporary Protected Status in the United States to eligible aliens who are nationals of a foreign state designated by the Attorney General, or who have no nationality and who last habitually resided in that state. The Attorney General may designate a state upon finding that the state is experiencing ongoing armed conflict, environmental disaster, or certain other extraordinary and temporary conditions that prevent nationals or residents of the country from returning in safety.

Effective on June 7, 1994, the Attorney General designated Rwanda for Temporary Protected Status for a period of 12 months, 59 FR 29440. This notice extends the designation of Rwanda under the Temporary Protected Status program for an additional 12 months, in accordance with sections 244A(b)(3) (A) and (C) of the Act.

This notice also describes the procedures with which eligible aliens who are nationals of Rwanda, or who have no nationality and who last habitually resided in Rwanda, must comply in applying for continuation of Temporary Protected Status.

In addition to timely re-registrations and late re-registration authorized by this notice's extension of Rwanda's Temporary Protected Status designation, late initial registrations are possible for some Rwandans under 8 CFR 240.2(f)(2). Such late initial registrants must have been ''continuously physically present'' in the United States since June 7,1994, and must have had a valid immigrant or non-immigrant status during the original registration period.

An application for Employment Authorization, Form I–765, must always